Applicants: Stewart Shuman et al.

Serial No.: Not Yet Known

Filed: Herewith

Page 3

## Amendments to the claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

## Listing of Claims:

- 1. (original) A method of covalently joining a DNA strand to an RNA strand comprising:
  - a. forming a topoisomerase-DNA intermediate by incubating a DNA cleavage substrate comprising a topoisomerase cleavage site with a topoisomerase specific for that site, wherein the topoisomerase-DNA intermediate has one or more 5' single-stranded tails; and
  - b. adding to the topoisomerase-DNA intermediate an acceptor RNA strand complementary to the 5' singlestrand tail under conditions permitting a ligation of the covalently bound DNA strand of the topoisomerase-DNA intermediate to the RNA acceptor strand and dissociation of the topoisomerase, thereby covalently joining the DNA strand to the RNA strand.
- 2-18. (canceled)
- 19. (original) A DNA-RNA molecule covalently joined by the method of claim 1.
- 20-22. (canceled)
- 23. (original) A covalently joined DNA-RNA molecule having a labeled 5' end.
- 24. (canceled)
- 25. (canceled)
- 26. (original) A method of tagging a 5' end of an RNA molecule comprising:
  - a. forming a topoisomerase-DNA intermediate by incubating a DNA cleavage substrate comprising a topoisomerase cleavage site with a topoisomerase specific for that site, wherein the topoisomerase-

Applicants: Stewart Shuman et al.

Serial No.: Not Yet Known

Filed: Herewith

Page 4

DNA intermediate has one or more 5' single-stranded tails; and

b. adding to the topoisomerase-DNA intermediate a 5'hydroxyl terminated RNA molecule complementary to the 5' single-strand tail under conditions permitting a ligation of the covalently bound DNA strand of the topoisomerase-DNA intermediate to the RNA molecule and dissociation of the topoisomerase, thereby forming a 5' end tagged DNA-RNA ligation product.

## 27-44. (canceled)

- 45. (original) A method of obtaining full-length gene sequences comprising:
  - a. isolating full-length mRNA;
  - b. attaching a DNA tag sequence to the isolated mRNA; and
  - c. synthesizing cDNA using the tagged mRNA as a template.

## 46-78. (canceled)

- 79. (original) A method of obtaining full-length gene sequences comprising:
  - a. isolating full-length mRNA by employing an affinity purification material;
  - b. decapping and dephosphorylating the isolated mRNA;
  - c. attaching a DNA tag sequence to the decapped, dephosphorylated mRNA, wherein the tag sequence comprises the sequence shown in Figure 11 and is attached by vaccinia DNA topoisomerase;
  - d. synthesizing cDNA using the tagged mRNA as a template;
  - e. amplifying the synthesized cDNA, wherein the amplification primers comprise an anti-coding sequence of the tag sequence (5') and a gene specific sequence (3'); and
  - f. inserting the amplified cDNA into an expression vector.